

AUTHORS: Barshcheva, A.S. and Gnuhev, V. S., Engineers
TITLE: High speed thin layer cyaniding during high frequency heating. ^{129-7-12/16}
(Skorostnoye tonkosloynoye tsianirovaniye pri nagreve
tokami vysokoy chastoty).
PERIODICAL: "Metallovedenie i Obrabotka Metallov" (Metallurgy and
Metal Treatment), 1957, No.7, pp.48-50 (U.S.S.R.)
ABSTRACT: The current type cyaniding in baths consisting of molten cyanide salts enables the saturation of the surface of components with carbon and nitrogen to depths of 0.1 to 0.6 mm and such components can then be hardened to $R_c = 60$. This process is impracticable for small components used in the instrument industry and the aim of the authors was to develop a process which permits obtaining a cyanide layer of a depth less than 0.1 mm with hardnesses of $R_c = 60$ and more. In this paper the results are described which were obtained by cyaniding by means of high frequency heating; two variants of this process were tried, namely, applying a paste prior to heating and cyaniding in molten potassium ferrocyanide. After degreasing the component with benzene a 3 to 4 mm thick layer of the paste was applied, on the top of which potassium ferrocyanide powder was sprayed and following that, the components were dried

Card 1/3

High speed thin layer cyaniding during high frequency heating. (Cont.)

129-7-12/16

for 40 to 60 minutes at 60 to 70 C. The heating was effected by a current of 200 kc/s, 160 kW. The thickness of the saturated layers varied between 0.023 and 0.07 mm. In the other variant potassium ferrocyanide (90%), barium chloride (10%), were placed in a crucible and molten by placing it into the first position of the inductor, then the components were immersed into the melt and heated in the second position of the inductor to 840-880 C. In both variants diffusion layer thicknesses of 0.04 to 0.06 mm were obtained and the hardness at the surface varied between $R_c = 59$ and 62. Using for such components medium carbon steels ensures obtaining a strong base and eliminates the possibility of squeezing through the thin diffusion layer during normal operation. The first variant requires a number of preparatory operations concerned with the manufacture of the paste and with drying of the paste coated components and, therefore, it is not recommended for series production. In the case of cyaniding in molten salts it is possible to obtain a given layer thickness with an accuracy up to 0.01 mm and the process can be fitted into an automatic

Card 2/3

BARSHCHEVA, A.S., inzh.

Annealing complicated machine parts with high-frequency currents.
Izobr. i rats. 3 no.5:9-12 My '58. (MIRA 11:9)
(Induction heating) (Metals--Heat treatment)

ZAZUBIN, A.I.; BARSHCHEVSKAYA, A.N.

Studying the distribution of gallium during the processing of
high-silicon bauxites by the hydrochemical alkali method. Trudy
Inst. met. i obog. AN Kazakh. SSR 9:103-105 '64. (MIA 17:9)

ZAZUBIN, A.I.; RASHCHEVSKAYA, A.N.

Investigating the distribution of silicon in the process-and-sintering
to th flow sheet of the Bayer-sintering processing of high-silicon
bauxites. Trudy Inst.met.i obog. AN Kazakh.SSR 11:22-23 '64.
(MIKA 18:4)

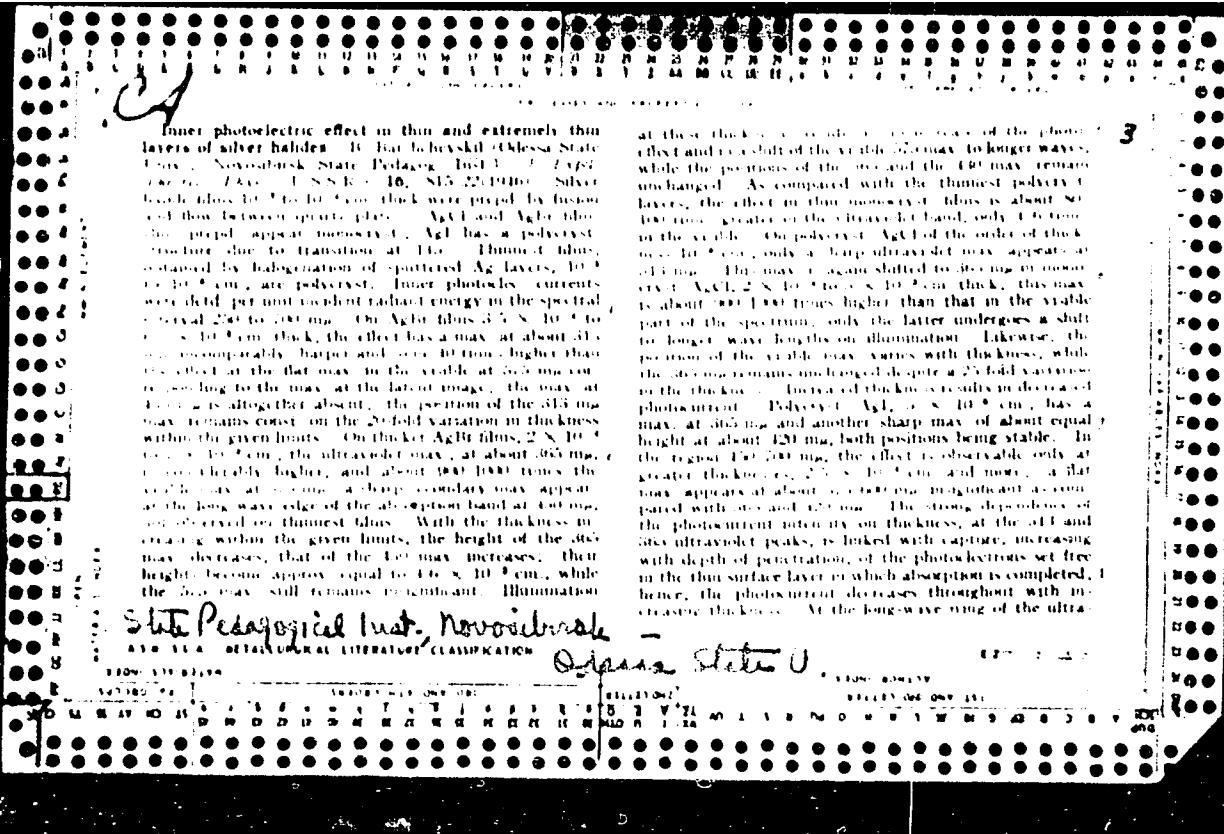
TARASHEV, V. S.; G. I. KUZNETSOV, T. N.

This will be distributed to the appropriate personnel for action
for adjustment by the Bureau of Counterintelligence, Defense Intelligence Agency,
and the FBI.

(MIA 18:16)

BARSHEVSKY, S.

"On the Importance of the Thickness of the Layer in the Inner Photoeffect,"
Zhur. Ekspер. i Teoret. Fiz., 12, Nos. 5-6, 1942
Inst. Physics, Odessa State U.



violet absorption band, the absorption could be much less than in its middle, electrons are set free also at some distance from the surface, and diffusion and capture play a relatively smaller role, hence the observed growth with increasing thickness of the violet zone at the edge of the ultraviolet absorption band. In thinner i.e., polycrystalline, the photocurrent drops as a result of illumination, owing to decompose of Ag halide and evaporation of source of photoelectrons; this effect loses its significance in thicker films where illumination gives rise to enhanced photoemission. Formation of colloidal Ag particles on illumination very likely underlies the observed increase and spectral shift to longer waves of the photocurrent may in the yellow-green region. N. They

BONCHOVSKIY, B.

FA 24T6

USSR /Chemistry - Photoelectric Effect
Chemistry - Cells, Photoelectric

Sep 1947
"Abnormal Inner Photoelectric Effect," B. Borsh-
chevskiy, 3 pp

"Zhur Fiziches Khim" Vol XXI, No 9 1947 - 1948

An abnormal (negative) inner photo effect results in lessening the electric conductivity of photo-semiconductors under the influence of light. The author observed this negative photo effect to study some photoelectric properties of silver halide salts, in poly-crystalline as well as macro-crystalline layers. The author, in experiments conducted at the Research and Investigation Cine-photo Institute, Laboratory of Photo Processing, Moscow, determined that the

24T6

negative inner photo effect is a result of secondary polarization processes.

24T6

USSR /Physics
Spectra, Ultraviolet
Photoelectric Cells

Sep 48

"Spectral Distribution of the Photoeffect in an
Antimony-Cesium Photoelement in the 2,483-5,770 Å
Region," B. A. Barshnevskiy, Sci Res Kino-Photo Inst
Moscow, 12 pp

"Zhur Tekh Fiz" Vol XVIII, No 9

Antimony-cesium photoelements sensitive to ultraviolet
light have been made in USSR for some time, although
??/49T10

USSR /Physics (Contd)

Sep 48

output is insufficient. Sensitivity to ultraviolet
light is obtained by using photosensitive antimony-
cesium layer, and also by using very thin special
ultraviolet glass. Describes experiment to determine spec-
trum characteristic of cell. Prints and discusses
results. Submitted 11 Mar 48.

32/49T102

BARSHNEVSKIY, B. A.

USSR/Physics

Light - Absorption
Films - Reflectivity

Mar 49

"The Absorption of Light in Dense Layers of Silver Halides," B. Barshchevskiy, Sci Res "Kinophoto" Inst, 4 pp

"Dok Ak Nauk SSSR" Vol LXV, No 1

Finds peculiarities in light absorption by thin silver-halide films which serve to clarify phenomena of the formation of a hidden photographic image, mainly on the surface of silver-bromide microcrystals in photographic emulsions, and also fact that the quantum yield determined by the photoelectric method is considerably less than unity (in AgBr, η equals 0.17 for λ equal to 254 m μ). Submitted by Acad S. I. Vavilov, 14 Dec 48.

29/49T103

(A)

Absorption of light by silver halides. B. A. Barshevskii (All-Union Kinophoto-Inst.). *Zhur. Ekspd. Teor. fiz.* 20, 117-20 (1950). - Absorption of light by thin layers of AgI_{tr} and AgI does not follow Lambert's law $I = I_0 e^{-kd}$, the coeff. k varying with the thickness d . With a given sample, e.g., polycryst. AgBr 1.4 $\times 10^{-4}$ cm. thick, the absorption $I = I_0$, where I_0 = transmission, becomes greater over the whole range of wave lengths when, after a series of measurements from 378 to 254 m μ , it is remeasured in the reverse order of wave lengths; the effect, which is more pronounced in longer waves, but also distinct in the near ultraviolet, is due to the formation of colloidal Ag during the 1st series of illuminations. The reflectance of polycryst. AgI_{tr} layers was found to vary from 10% in 378 to 14% in 254 m μ ; for polycryst. AgBr it was found to be 15.5% in 378 m μ , 16.5% in 435, and 14% in 254. For monocryst. AgI_{tr} (produced between 2 quartz plates), the reflectance varied from 10% in 378 to 34% in 254, and for monocryst. AgCl it was on the avg. 10% over the whole range. I_0 (calcd. with the reflected light deducted) varies surprisingly little with the thickness of the layer, 2.2 = 42 $\times 10^{-4}$ cm. for polycryst. AgBr, 3.2 = 53 $\times 10^{-4}$ cm. for polycryst. AgI, and 2.45 = 30.8 $\times 10^{-4}$ cm. for monocryst. AgBr. Thus, in 365 m μ , a 2.2 $\times 10^{-4}$ cm. 1 layer of AgI_{tr} absorbs 81%, a 3.2 layer of AgI, 70% of the unreflected light, with 20 times thicker layers the absorption increases only 9 and 2%, resp. Thinnest layers show a strikingly strong absorption. The effect of the structure of a particular layer, grain size and nature of the contact between the grains, on the absorption is illustrated by the difference between a 15 $\times 10^{-4}$ and a 20.3 $\times 10^{-4}$ cm. film of AgI_{tr}; the 1st absorbs strongly over the whole spectrum, the 2nd is transparent as far as

the ultraviolet. In the range 378-310 m μ , fluctuations of the absorption from one layer to another are greater than in the absorption band; in this range, evidently, absorption is determined mainly by the granular structure, and is markedly weaker in a monocryst. film. In the range 435-405 m μ , polycryst. and monocryst. AgI_{tr} absorb approx. to the same degree, but in the ultraviolet monocryst. AgI_{tr} absorbs much more strongly. In AgCl, absorption in 405 m μ is about the same as in 435-378, and much weaker than in the same range in AgI_{tr}; in 365 m μ , AgCl, 6.8 $\times 10^{-4}$ cm., transmits 47.2% as against only 4% for AgI_{tr} 7.62 $\times 10^{-4}$ cm. The Lambert-law coeff. k vary strongly with the thickness and are consequently meaningless. This inapplicability of the law is clearly due to the strong absorption which takes place in the very thin surface layer. An absorption coeff. can be conveniently defined by $k' = A = (I_0 - I)/L$, indicated by extrapolation to a thin layer of the order of 10⁻⁴ cm. The conclusion that Ag halides absorb light primarily in the outer surface layer is in accord with the decrease of the photoconductivity with increasing thickness. The strong surface absorption is due to an internal photoelectric effect, whereby the absorbing electron is removed from the lattice point. These peculiarities of the absorption account for the known preferential formation of the latent photographic image at the surface of the AgI_{tr} grains, and for the fact that the quantum efficiency is substantially less than unity.

N. Thom

168T97
BARSHCHEVSKIY,

USSR/Physics - Photoeffect, Internal
Absorption, Light

Sep 50

"Relation Between the Internal Photoeffect and
the Absorption of Light in Silver Halide Salts,"
B. Barshchevskiy, All-Union Sci Res Kinophoto
Inst

"Zhur Eksper i Teoret Fiz" Vol XX, No 9, pp 795-
806

Studies spectral distribution of internal photo-
effect for equal absorption of energy in AgBr and
AgCl. Shows there are regions of selectivity of in-
ternal photoeffect. Increase of photoeffect in

USSR/Physics - Photoeffect, Internal
(Contd)

168T97
Sep 50

"Absorption of Light, Internal Photoeffect, and Photo-
graphic Sensitivity in AgI," B. Barshchevskiy, All-
Union Sci Res Kinophoto Inst

monocrystallic salts is much greater, as result of in-
stantaneous illumination, than corresponding increase
of absorption in them. Submitted 25 Feb 50.

PPROVED FOR RELEASE: 06/06/2000 CIA-RDP86-00513R000203710015-2
"Zhur Eksper i Teoret Fiz" Vol XX, No 9, pp 862
Absorption A of light in

168T97

BARSHCHEVSKIY, B.

16ST104

USSR/Physics - Photoeffect, Internal
Sensitivity, Photographic Sep 50

"Absorption of Light, Internal Photoeffect, and Photo-
graphic Sensitivity in AgI," B. Barshchevskiy, All-
Union Sci Res Kinophoto Inst

"Zhur Eksper i Teoret Fiz" Vol XX, No 9, pp 862

Letter to editor considers absorption A of light in
AgI layer $2/10^5$ cm thick as function of wave length
L (A varies roughly from 100 to 20% as L goes from
250 to 650 millimicrons); also photoeffect vs wave
length according to incident and absorbed energies.
Submitted 25 Mar 50.

SA

A 53

537.221

7829. Image formation by electrographic means. D. Barshchovskii and V. Lavrouchik. Letter in J. Tech. Phys., USSR, 20, 1273-4 (Oct., 1950) In Russian.

A latent image of a conductor may be formed on an insulator such as S or Arapex (not glass or ebonite) if they are placed in contact and the conductor is charged to \pm 5 kV. The image may be made visible by dusting with powdered resin or lycopodium powder; if resin is used, the image may be "fixed" by gentle warming. J. C. E. Jennings

DARSHCHEVSKII, S.

"Internal Photo-Effect and Light Absorption by Silver Halides," Usp.
nauch. fot., No.1, 1951

BARSHEVSKIY, B.

3

Energy Spectrum of the Electrons in Silver Bromide Crystals. B. BARSHEVSKIY.
Zh. Exper. Teor. Fiz. 1952, 22, 511-512.—The photoelectrons required for the formation of the latent image are torn from the halide ions of the crystal lattice of the silver halide by light radiation of a wavelength inside the characteristic absorption band. If an electric field is applied to the crystal during the latent image formation, a photocurrent is set up, i.e., the photoelectrons, before being separated at the centres of recombination, are transferred to conductivity bands of the crystal. During the photoconductive process the electrons may reach the positive electrode or be captured, thus eventually forming centres or sub-centres of a latent image. Using experimental data on the spectral distribution of the photo-effect, the author presents a schematic diagram of the electron

BARSHEVSKIY, B.N.

Modulus of deformation and the Poisson coefficient for an incoherent soil. Trudy LIEI no.57:42-49 '65.

Determining stresses in soil with variable characteristics of deformability. Ibid.:50-68

Contact stresses along a foundation base supported by soil with varying depth deformability characteristics. Ibid.:69-88

Strength calculation of rigid retaining works erected on an incoherent soil. Ibid.:89-97 (MIRA 18:8)

BARSHCHEVSKIY, B.U., dotsent, kand. fiziko-matem. nauk; KUROVA, A.V.,
red.; KLEYMAN, L.G., tekhn. red.

[Physics; dispersion of light] Fizika; dispersiia sveta. Konspekt
lektsii dlja studentov II kursa vsekh spetsial'nostei. Moskva,
Vses. zaochnyi institut inzhenerov zhel-dor. transp., 1960. 13 p.
(MIRA 14:7)

(Refraction)

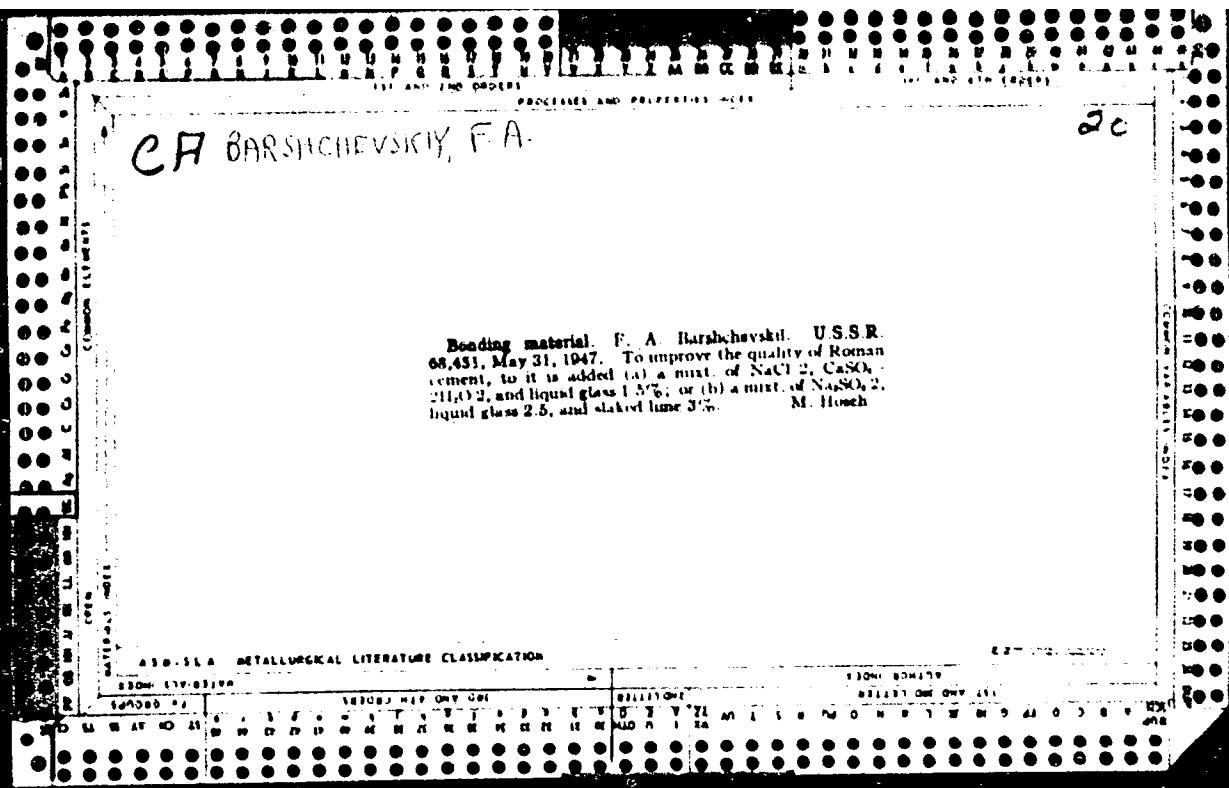
BARSHEVSKIY, B.U., dotsent, kand. fiz.-mat. nauk; KHONDKARYAN, N.G., red.;
KLEYMAN, L.G., tekhn. red.

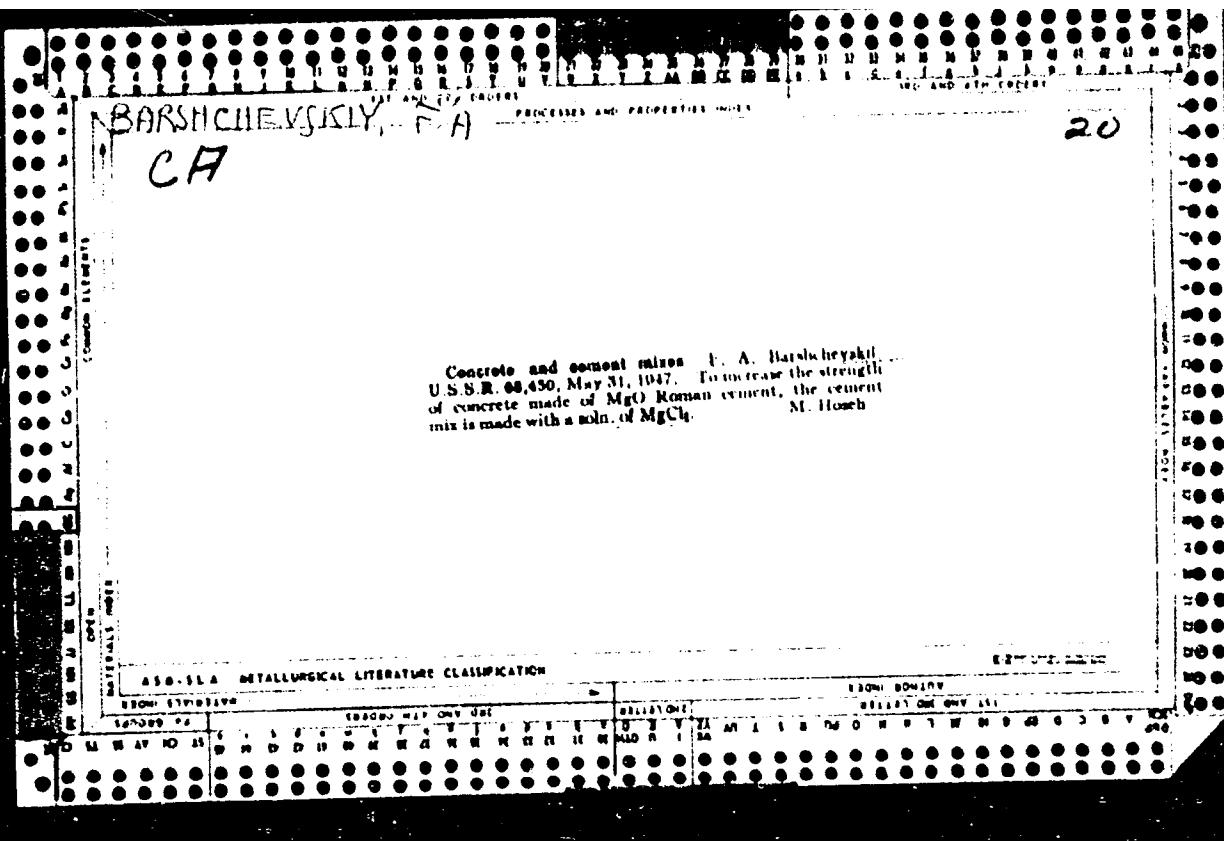
[Electrostatics; abstract of lectures of the second course of all
faculties] Elektrostatika; konспект lektsii dlja studentov II kursa
vsekh spetsial'nostei. Moskva, Vses. zaochnyi in-t inzhenerov zhel-
dor. transp., 1960. 31 p. (MIRA 14:7)
(Electrostatics)

Br. 96, BARSHCHEVSKIY, F. A.

BI-1. Chemical Engineering,
Plant, Machinery.

Improving the efficiency of tunnel dryers. F. A. Barshchhevskiy
(Sov. Krem. Prom., 1947, No. 6, 14; Byul. Czass., 46; 1948,
218a) — Two methods are proposed. One consists of linen baffles
hung from the roof across the tunnel between every other wagon;
these are pushed up and fall into place again as the wagons pass.
The other consists of metal dampers which are raised and lowered
as required from outside. The purpose of both devices is to force
the gases along the tunnel in zigzag manner from top to bottom
of the tunnel instead of travelling in a straight line along the top.
R. H. Crase





15-57-3-3391

Translation from: Referativnyy zhurnal, Geologiya, 1957, Nr 3,
p 135 (USSR)

AUTHOR: Parshchevskiy, F.

TITLE: The Classification of Brick and Tile Clays by the Composition of Their Grains (Klassifikatsiya kirkpichno-cherpichnykh glin po zernovomu sostavu)

PERIODICAL: Stroit. materialy, izdeliya i konstruktsii, 1956, Nr 7,
p 30

ABSTRACT: The author recommends a new classification of brick and tile clays by the composition of their grains. The clays are divided into three subgroups: sandy, clay proper, and powdery. The content of the three principal fractions for each subgroup is given in the table.

Card 1/2

The Classification of Brick and Tile Clays (Cont.)

18-57-3-3391

| Name of clay according to new classifi- cation | Content of fraction, percent | | |
|---|------------------------------|-------|--------|
| | sand | clay | powder |
| Sandy clay | 40-70 | 30-60 | 30-60 |
| Clay | 20-10 | 30-60 | 30-60 |
| Powdery clay | 0-20 | 30-60 | 40-70 |

Card 2/2

S. P. Sh.

BARSHCHEVSKIY, F., kandidat tekhnicheskikh nauk.

Determining the amount of water to be used in normal slaking
of clay materials. Stroi. mat., izdel. i konstr. 2 no.8:27
Ag '56. (MLRA 9:10)

(Clay)

RASHCHEVSKIY F.

RASHCHEVSKIY, F., kand.tekhn.nauk (Kiyev)

Determining the suitability of clayey deposits for use in
brickmaking. Stroi.mat. 3 no.11:30-31 N '57. (MIRA 10:12)
(Clay--Testing)

BARSHCHEVSKII, F. [Barshchev's'kyi, F.], kand.tekhn.nauk

Making Parker's cement in brick factories. Sil', bud. 7 no.4:
11-12 Ap '57. (MIRA 12:11)
(Ukraine--Cement)

RARSHCHEVSKIY, F. [Burshchevs'kiy, F.], kand.tekhn.nauk

Classifying raw materials and batching in making bricks. Sil'.bud.
8 no.2:12-13 F '58. (MIRA 13:7)
(Brickmaking)

LYSIN, B.S., akademik; BARSHCHEVSKIY, F.A. [Barshchevs'kyi, F.A.]

Studying the physical and chemical properties of Dnieper marl as a raw material for the production of local adhesives. Dop.AN URSR no.9:1258-1262 '60. (MIRA 13:10)

1. Kiyevskiy politekhnicheskiy institut. 2. AN USSR (for Lysin).
(Marl) (Cement)

BARSHCHEVSKIY, G., kand. tekhn. nauk

Performance of synchronous generators without excitation
in electric ship systems. Mor. flot 23 no.8:22-25 Ag '63.
(MIRA 16:11)

RUMYACHEVSKIY, G.G., kand. tekhn. nauk, dozent. (Pushkin)

Electromagnetic moment of a reactive machine. Elektricheskiye
no.3:74-78 Mr '65. (MIRA 18:6)

NAZHOR V. N.,

"Concerning the Nature of Activ. Centers in Photocatalytic Reactions." Card
Chem Sci, Moscow Order of Lenin State U imeni M. V. Lomonosov, 17 Dec 54.
(U, 7 Dec 54)

Survey of Scientific and Technical Dissertations Defense'd at US & Higher
Educational Institutions (12)
SC: Sub. No. 556, 24 Jun 55

Barshchevskiy, I.N.

USSR/Chemistry - Physical chemistry

Card 1/1 Pub. 147 - 11/27

Authors : Nikolayev, L.A., and Barshchevskiy, I.N.

Title : Photocatalytic properties of ZnO in methylene blue reduction reactions. I.

Periodical : Zhur. fiz. khim. 28/2, 265-270, Feb 1954

Abstract : It was established experimentally that ZnO in the role of a catalyst strongly accelerates the reduction of methylene blue with formaldehyde under the effect of light. Pure ZnO was found to be less active than ordinary commercial compounds. An active photocatalyst was obtained by adding Mn and Cu compounds to the pure oxide. Cu appears to be a stronger activating agent than Mn. The Cu-ions by themselves showed no catalytic effect on the reaction between methylene blue and formaldehyde. The basic kinetic laws, governing the reaction of photo-reduction of dyes in the presence of ZnO as catalyst, are presented. Six references : 4-USSR; 1-USA and 1-German (1926-1948). Table; graphs.

Institution : The I.V. Stalin Institute of Transportation Engineers, Moscow

Submitted : April 21, 1953

Barshchevskiy, I. N.

USSR/Chemistry - Physical chemistry

Card 1/1 Pub. 147 - 12/27

Authors : Nikolayev, L.A.; and Barshchevskiy, I.N.

Title : Photocatalytic properties of ZnO in methylene blue reduction reaction. II.

Periodical : Zhur. fiz. khim. 28/2, 271-274, Feb 1954

Abstract : The effect of Cu-ion concentrations in ZnO on the photocatalytic activity of the latter in the reaction between formaldehyde and methylene blue was investigated. It was found that the catalytic activity, relative to 1% of Cu (specific activity), increases sharply in ratio to the decrease of the total Cu concentration in ZnO. A direct proportionality between the amount of Cu introduced and the photocatalytic activity was observed at greater Cu concentrations. A sharp increase in specific activity was seen in the zone of Cu concentration in which the luminescence nature changes considerably. Pb-ions appeared to be much stronger activators of the photocatalytic activity of ZnO. The luminescence of ZnO samples containing Pb changes with the reduction in Pb content. Two USSR references (1952 and 1954). Tables; graphs.

Institution : The I.V. Stalin Institute of Transportation Engineers, Moscow

Submitted : April 21, 1953

BARSHCHEVSKIY, I.N.

USSR/Chemistry - Photocatalysts

Card 1/1 : Pub. 147 - 17/27

Authors : Nikolaev, L. A., and Barshchevskiy, I. N.

Title : Photocatalytic properties of ZnO in the reduction reaction of methylene blue. Part 3. Poisoning of photocatalysts

Periodical : Zhur. fiz. khim. 28/12, 2211-2214, Dec 1954

Abstract : Experiments showed that the ions of heavy metals poison the ZnO photocatalyst but do not affect its luminescence, whereas nitrate and nitrite ions poison the photocatalyst and extinguish its luminescence. Substances which extinguish the luminescence of a photocatalyst must, of course, also be catalytic poisons since they distort one of the component parts of the active center. It is assumed that the active center of a photocatalyst has a double (polar) structure. One of the component parts of the catalyst center is the luminescent center the other is probably the distorted part of the lattice of the basic substance (ZnO) oriented in close vicinity to the luminescence center. Three USSR references (1950 and 1954). Tables; graphs.

Institution : Institute of Transportation Engineers, Moscow

Submitted : May 3, 1954

AF701597

TREASURE ISLAND BOOK REVIEW

AID 798 - S

BARSKHEVSKIY, I. N. and L. A. NIKOLAYEV. (Moscow Institute of Railroad Engineers).

OSOBNOSTI STROYENIYA AKTIVNYKH TSENTROV V FOTOKATALIZE (Characteristics of formation of active centers in photocatalysis). In Problemy kinetiki i kataliza (Problems of Kinetics and Catalysis), vol. 8. Izdatel'stvo Akademii Nauk SSSR, 1955. Section I: Effect of light on the adsorbability of solids. p. 61-67.

The activity of photocatalysts depends greatly on the presence of foreign agents. Bi^{++} , Pb^{++} , Tl^{+++} , Ag^+ , Cu^+ and Mn^{++} introduced into the lattice of ZnO greatly increase the photocatalytical activity of the latter. A catalyst containing Pb-ions is 5 times as active as a catalyst containing Cu-ions. Ions of Ba, Ca, Cu and Pb, gaseous H_2S as well as nitrite and nitrate ions, poison photocatalysts.

To establish the dependence of the catalytic activity on luminescence, the effects of a decrease in the concentration of the activator (Cu and Pb) on the luminescence has been studied and data presented in tables. Since positive and negative ions may serve as catalyst poisons, and some substances

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BARSHCHEVSKIY, I. N. and L. A. NIKOLAYEV. Osobennosti . . . AID 796 - 3

may inhibit catalysis without affecting luminescence, it is assumed that the active center has a bipolar structure. Luminescence centers are a part of active catalytic centers, for example, ions of copper or lead are centers of luminescence and at the same time parts of bipolar catalytic centers. Substances inhibiting catalysis may not quench the luminescence. Substances which quench the luminescence are catalytic poisons.

ZnO activated by ions of heavy metals (Bi, Pb, Tl, Ag, Cu, and Mn) shows high photocatalytic activity. These ions are a part of the active center which are assumed to be bipolar. The active centers of photocatalysts have some characteristics which active catalytic centers do not possess. Calcination of ZnO (activated by Pb) over a period of 2 hours at 600°C decreases the activity by 50% at 800°C - by 90%. Compression at 4000 kg/cm² decreases the activity by 50%, at 10,000 kg/cm² by 80%.

Various methods of activating zinc oxide are discussed. Other semiconductors, such as titanium dioxide, cadmium oxide, lead oxide and cupric oxide, were also investigated. Introduction of ions which activate zinc oxide did not affect the above-mentioned semiconductors.

2/3

YARSHCHEVSKIY, I. N. RUD'YA M. NIKOLAEV. Obshchennosti ... AID 798 - S

The systems: formaldehyde-thionine, glycerin-methylene blue, glycern-thionine in the presence of ZnO have been investigated. ZnO proved to be an active catalyst.

BARSHCHEVSKIY, I.N.

The nature of the composition of active centers in photo-catalysts. I. N. Barshchevskiy and L. A. Nikolayev (Inst. of Transport, Eng., Moscow). Problemy Khimii i Kataliz, Akad. Nauk S.S.R., 8, 61-7 (9155).—ZnO, after activation by ions of heavy metals, appears to be a specific photocatalyst for reduction of methylene blue (I) and of thionine by CH_3O , glycerides, or succinic acid. Ions of such heavy metals as Cu^{++} , Pb^{++} , Bi^{+++} , Tl^{+++} , Ag^+ , Zr^{4+} , Mn^{++} , enter the photocatalyst in the state of active centers and generate a specific distribution of energy levels, which favors photocatalysis. The active centers are liable to formations with irregular structures that are not in equilibrium with their surroundings. They have a double structure: one portion serves for light absorption, the other for catalysis. The surface of such a photocatalyst has 3 zones: one where the ZnO shows the regular lattice, one where the ions of the activator are located, and one that surrounds these defects. The concn. of the heavy metals is crit.; thus, changing the amt. of Cu from 0.0255% to 0.00087% in slow steps changes the color of the luminescence from green to weak green to yellowish green to yellowish brown to yellow-orange to red-orange, and similarly changing the amt. of Pb from 0.0026 to 0.00187% changes the luminescence from yellow-brown to red. The following materials have an extinguishing effect on the luminescence: KI, KBr, KCNS, K_2CO_3 , $\text{K}_2\text{S}_2\text{O}_8$, KClO_4 , KBrO_3 , KIO_4 , KMnO_4 , NaClO_4 , NaHPO_4 , NaSCN , Na_3F , $\text{Na}_2\text{S}\cdot\text{H}_2\text{O}$, HCO_3Na , NaCl , La acetate, $\text{BaCl}_2\text{H}_2\text{O}$, $\text{CaCl}_2\cdot\text{H}_2\text{O}$, Pb(OAc)_2 , CoCl_2 , MnSO_4 , $\text{CICH}_2\text{CO}_2\text{H}$, BzOH , pyrogallol, rosolic, oxalic, sulfanilic, formic, adipic, and oleic acids, phenol, water, I, the leucobase of I in the presence of other reduction products of I, CH_3O , methyl violet, hydrazine, hydroquinone, EtOH , and diphenylamine. Werner Jacobson

BARSH CHEVSKIY, I. N.
USSR/ Chemistry - Photo-catalysis

Card 1/2 Pub. 22 - 31/50

Authors : Nikolayev, L. A., and Barshchevskiy, I. N.

Title : Catalytic properties of sensitizers

Periodical : Dok. AN SSSR 100/1, 119-122, Jan. 1, 1955

Abstract : The nature of active centers on the surface of ZnO is explained from the view point of photo-catalysis. The question on whether electron levels of admixtures are of importance for the photo-catalytic effect and whether catalysis centers are in approximate ratio to luminescence centers is debated. It was found that pure ZnO free of heavy metal admixtures is not a photo-catalyst and does not accelerate the dye reduction process when exposed

Institution : The I. V. Stalin Institute of Railroad Engineers, Moscow

Presented by : Academician A. N. Terenin, June 26, 1954

Periodical : Dok. AN SSSR 100/1, 119-122, Jan. 1, 1955

Card 2/2 : Pub. 22 - 31/50

Abstract : to light. ZnO containing an admixture of heavy metal ions was found to be a highly active photo-catalyst. H₂S was established to be a catalytic poison for photo-catalysts containing Pb and Cu. The effect of poisoning of photo-catalysts with Ca. and Ba ions does not result in a change in the luminescence. Ten USSR references (1940-1954). Graphs.

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SOV/65-4-3-26/31

AUTHOR: Barshchhevskiy, I.N.

TITLE: The Poisoning of Photo-Catalysts by Copper and Lead Ions

PERIODICAL: Khimicheskaya nauka i promyshlennost', 1959, Vol 4, Nr 3, pp 412-413 (USSR)

ABSTRACT: Zinc oxide which is used as photo-catalyst in the preparation of leuco-methylene blue and leucothionine is poisoned due to active centers on its surface. The structure and the properties of these centers are studied here. Formaldehyde is used as electron donor and sulfate of bivalent copper and acetate of bivalent lead as poisons. The reaction is carried out in a calibrated glass photoreactor. The activity of the reactor depends on the equilibrium concentration of the poison ions. The ions block the active centers. Lead ions show a stronger effect than copper ions. Kobozev's conception of the latent parameters of catalytic processes may be applied to the poisoning of photocatalysts [Ref 6]. There are: 1 graph and 6 Soviet references.

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The Polymerizing of Photo-Catalysts by Copper and Lead Ions

SOV/63-4-3-22/31

ASSOCIATION: Moskovskiy institut inzhenerov zheleznodorozhnogo transporta, kafedra
khimii (Moscow Institute of Railroad Transportation Engineers, Chair
of Chemistry)

SUBMITTED: January 14, 1959

Card 2/2

5 (4)

AUTHOR:

Barshchevskiy, I. N. (Moscow)

SOV/76-37-5-12/33

TITLE:

The Photocatalytic Properties of Zinc Oxide in the Reaction
of the Reduction of Methylene Blue (Fotokataliticheskiye
svoystva okisi tsinka v reaktsii vosstanovleniya metilenovoy
sini). 4. The Poisoning of Photocatalysts by Nitro Compounds
(4. Otravleniye fotokatalizatorov nitrosoyedineniyami)

PERIODICAL:

Zhurnal fizicheskoy khimii, 1959, Vol 33, Nr 5,
pp 1071-1079 (USSR)

ABSTRACT:

The relation between the molecular structure of the poisons and their toxic effect as well as the structure of the active centers of the photocatalyst were investigated. The methods used were already described in previous papers (Refs 1-3). Zinc oxide activated by lead ($\alpha = 3.754 \cdot 10^{-3} \%$ Pb²⁺) was investigated. The toxic effect of sodium nitrate, sodium nitrite, o-, m-, and p-nitrophenol was examined; moreover, the effect of dinitrophenols, thiopheneless benzene, phenol, and pyridine was investigated. Figures 1-4 show that the apparent activity initially decreases rapidly with increasing concentration of the poison but then approaches the axis of abscissas as a flat asymptote. By apparent activity, the

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The Photocatalytic Properties of Zinc Oxide in the
Reaction of the Reduction of Methylene Blue, 4. The Poisoning of
Photocatalysts by Nitro Compounds

author understands the photocatalytic activity which is not reduced to the unit of the concentration of the poison. Toxicity increases in the order $\text{NaNO}_3 < \text{NaNO}_2 < m\text{-NO}_2\text{C}_6\text{H}_4\text{OH} <$ $< p\text{-NO}_2\text{C}_6\text{H}_4\text{OH}$. o-Nitrophenol and the other compounds mentioned showed no toxic effect. The strong toxicity of p-nitrophenol corresponds to its high dipole moment whereas the ineffectiveness of o-nitrophenol might be caused by its considerably lower dipole moment. The anion being formed in a molecule with OH group probably is bound more or less tightly to the cation of the lattice of the photocatalyst. In order to achieve toxic effect, however, the NO_2 group and a sufficiently high dipole moment have to be present in addition to the OH group. According to the author, this dependence on the dipole moment proves that a great shift of the active center towards the periphery takes place. In addition, the poisoning hyperbolae, the form of which is similar to that of the poisoning of metallic catalysts in

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The Photocatalytic Properties of Zinc Oxide in the SCV/75-33-5-18/33
Reaction of the Reduction of Methylene Blue. 4. The Poisoning of
Photocatalysts by Nitro Compounds

"dark" catalysis, were calculated. Figures 5, 6, and 8 show the dependence of activity on the logarithm of the equilibrium concentration of the anion of the poison. The ratio between the inclination angle of the steep branch of the hyperbola and the inclination angle of the asymptotically flattened branch of the hyperbola are compared with the dipole moments. Gratitude is expressed to Z. M. Finogenova and G. A. Pevtsov for supplying the pure preparations and to A. M. Dlin for recommending the method and equations for the determination of the results by means of the methods of mathematical statistics. There are 8 figures and 11 references, 12 of which are Soviet.

ASSOCIATION: Institut inzhenerov zheleznodorozhnogo transporta im. I. V. Stalina, Moskva (Institute of Railroad Engineers imeni I. V. Stalin, Moscow)

SUBMITTED: October 21, 1957
Card 3/3

S/076/62/056/132/CN1/009
B101/B*02

AUTHOR: Barshchevskiy, I. N.

TITLE: Photocatalytic properties of zinc oxide in the reduction of methylene blue. V. Poisoning by metallic ions and some causes of poisoning and activity of photocatalysts

PERIODICAL: Zhurnal fizicheskoy khimii, v. 36, no. 2, 1962, 249 - 259

TEXT: ZnO , activated by Pb ions ($\alpha = 3.574 \cdot 10^{-3} \text{ g Pb}^{2+}$) was subjected to the action of formaldehyde (donor) and methylene blue (acceptor) in oxygen-free N_2 in the presence of metallic ions. The rate $1/\tau_{1/2}$ of the change in the color was measured with an $\Phi K-5$ ($\Phi K-5'$) photocolorimeter. The concentration c_p of the poisoning ions was between 0 and $24 \cdot 10^{-6} \text{ g-dm}^{-3}$. The results are: Li^+ , Sr^{2+} , Ca^{2+} , Pb^{2+} , Ce^{2+} , Ni^{2+} , Cu^{2+} , Al^{3+} have a poisoning effect; Zn^{2+} , Cd^{2+} , Ga^{3+} , In^{3+} , NH_4^+ , SO_4^{2-} and CH_3COO^- have no effect. At the beginning the poisoning curve is a hyperbola which then passes into a straight line (Fig. 7). The poisoning is caused by non-

Photocatalytic properties of zinc

S/076/62/056/002, 001, 006
B'01/B102

screened structure of the poisoning ions, and if poisoning is due to the screening effect also by the high dipole moments of the poison molecules the mobility constant's and the presence of OH \cdot Zn $^{2+}$, probably also Pb $^{2+}$ pass into solution from which poison ions or dye ions and formaldehyde molecules are adsorbed. The adsorption - desorption equilibrium is the main reason of the activity and the poisoning of photocatalysts. Coordinate bonds are formed between the poison ions and the active bipolar centers. The energy level (local impurity levels) of the activator ions in the ZnO lattice differs from that of the Zn ions or molecules in solution. The active bipolar centers of the ZnO activated by Cu $^{2+}$ or Pb $^{2+}$ (according to A. N. Terenin. Fotokhimiya krasiteley (Photochemistry of Dyes) Izd-vo AN SSSR M.-L., 1947) are poisoned by the same ions. Photocatalysis and photosynthesis are inhibited by Cu $^{2+}$ and Cr $^{2+}$. A similar course can be expected for the transmission of light energy in the active centers of photocatalysis and photosynthesis. The poisoning of ZnO consists of true transitional and latent poisoning. In the latter, bipolar centers produce no effects. The sections of the curve depicting the pH dependence of the

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Photocatalytic properties of zinc ...

S/076/62/036/002/001/009
B101/B102

equilibrium concentration of Cu⁺ in solution after adsorption on ZnO correspond to the branches of the poisoning curve. The maximum of this curve approximately corresponds to the concentration of the poison ions in solution at which true poisoning takes place. Beside bipolar centers also stable active centers caused by Frenkel' defects are present on the ZnO surface: the interstitial spaces contain superstoichiometric Zn-ions (or molecules) and the vacancies are located at the nodes. The activator and Zn ions (or molecules) in the interstices facilitate the formation of stable centers without taking part therein. These centers are not poisoned by metallic ions but are relatively stable against thermal and mechanical treatment. Both types of centers take part in photocatalysis. N. I. Kobozev (Zh. fiz. khimii, 33, 1002, 1959) is mentioned. There are 7 figures and 31 references: 25 Soviet-bloc and 6 non-Soviet-bloc.

ASSOCIATION: Moskovskiy institut inzhenerov transporta (Moscow Institute of Transportation Engineers)

SUBMITTED: March 17, 1960

Card 3/*8*

S/076/52/036/000 06/000
B152/4110

AUTHOR: Burshchevskiy, I. N.

TITLE: Photocatalytic properties of zinc oxide in the reduction of methylene blue. VI. The dependence of the activity of the photocatalyst on the ionization potentials of atoms

PERIODICAL: Zhurnal fizicheskoy khimii, v. 36, no. 2, 1962, 369 - 372

TEXT: The dependence of the activity of zinc oxide activated by lead ions in its lattice ($a = 3.754 \cdot 10^{-3} \text{ Pb}^{2+}$) on the ionization potential of poisoning metal ions has been investigated. For this purpose, specific activities, i. e., activity per g-ion/ml of poison in the solution were calculated. As an ion adsorbed on the surface of a semiconductor is coordinately bound to the electrons of activated zinc oxide, the activities were related to 1 ev of the first, second, or third ionization energy. The dependence of the specific activity of the poisoned photocatalyst on the ionization potential of the poisoning ions is approximately a hyperbola (Fig. 1):

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B152/B110

$a_i^* = k/I + b \text{ min}^{-1}/\text{g-ion/ml/ev}$ (1), where $k = 2.83 \cdot 10^5$ and $b = 0.06 \cdot 10^5$.
The specific activity of the photocatalyst poisoned by metal ions may be expressed by the sequence

$$\begin{aligned} a_{\text{Cs}^+}^* > a_{\text{Rb}^+}^* > a_{\text{K}^+}^* > a_{\text{Na}^+}^* > a_{\text{Li}^+}^* > a_{\text{Ti}^{4+}}^* > a_{\text{Ag}^+}^* > a_{\text{Ba}^{2+}}^* > a_{\text{Sr}^{2+}}^* > a_{\text{Ca}^{2+}}^* > a_{\text{Pb}^{2+}}^* > \\ > a_{\text{Mg}^{2+}}^* > a_{\text{Co}^{2+}}^* > a_{\text{Ni}^{2+}}^* > a_{\text{Cu}^{2+}}^* > a_{\text{Al}^{3+}}^* \end{aligned} \quad (3)$$

These and previous results suggest that if in a monomolecular layer at the surface of zinc oxide there are ions of metals, nitro compounds, or methylene blue, the electrons of the active bipolar center of the photocatalyst will be blocked in the Coulomb field of adsorbed ions. It is assumed that the poisoning of the active center is the stronger, the greater is the dipole moment or the ionization potential of the metal atom. In the limiting case, the electrons may emerge from the surface of the semiconductor, the adsorbed ion becomes an atom and remains in the monomolecular layer at the surface of the positively charged zinc oxide. The toxication of the metal ions and the activity of the zinc oxide are periodic functions of the atomic number of the elements (Fig. 2). By comparing the values of enthalpies ΔH_h , entropies ΔS_h , and free energies

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Photocatalytic properties of zinc...

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3152/B110

ΔE_h of hydration of metal ions in infinitely diluted solutions at 25°C with the sequence of activities, it is found that the blocking energy is the smaller and activity the greater, the smaller are the thermodynamic data. There are 2 figures and 15 references: 14 Soviet and 1 non-Soviet. The reference to the English-language publication reads as follows: Handb. of chem. & phys., 37-th Ed. Chem. Rub. Publ. Co., Cleveland, Ohio, 1955-56, 2341.

ASSOCIATION: Moskovskiy institut inzhenerov transporta (Moscow Institute of Transportation Engineers)

SUBMITTED: December 21, 1960

Fig. 1. Specific activity a_i^* of the photocatalyst as a function of the ionization potential I of the atoms. (1) experimental values; (2) calculated values.

Legend: (a) $\text{min}^{-1}/\text{g-ion}\cdot\text{ml}^{-1}\cdot\text{ev}^{-1}$.

Card 3/4

BARSHCHEVSKIY, I.N., dots.

[Physicochemical properties of semiconductors] Fiziko-khimicheskie svoistva poluprovodnikov; uchetnoe posobie dlja studentov starshikh kursov MIIT-a. Moskva, Mosk. in-t inzhenerov transporta, 1962. 60 p. (MIRA 16:4)
(Semiconductors--Handbooks, manuals, etc.)
(Transistors--Handbooks, manuals, etc.)

BIRKOVSKIY, I.N.

Photocatalytic properties of antimony trioxide in the
reduction of methylene blue. Part 1. Zhur. fiz. khim.
no.11:2515-2517 N°62.

(MIRA 17:5)

1. Moskovskiy institut inzhenerov zheleznychodozhnego transporta.

MITYUREV, A.K.; BARSHCHEVSKIY, M.M.

Effect of shale coarseness on operation indexes and lining wear of compartment kilns in the Kohtla-Jarve shale reprocessing combine.
Trudy VNIIPS no.3:183-190 '55. (MLRA 8:12)
(Baltic Sea region--Oil shales) (Hydrocarbons)

MITYUREV, A.K.; BARSHCHEVSKIY, M.M.

Lowering the dust content of producer tar. Trudy VNIIPS no.4:
190-198 '55.
(Slantsy--Gas producers) (Dust--Removal) (MIRA 13:4)
(Tar)

BARSHCHERVSkiY, M.M.; MITYUREV, A.K.

Care of refractory material and major repairs in industrial compartment kilns. Trudy VNIIPS no.5:154-171 '56. (MLRA 10:5)
(Oil shales--Refining)

BARBIEU-VERA, N. M.

APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000203710015-2"

BARSHCHEVSKIY, M.M.; BEZMOZGIN, E.S.; ZAGLODIN, L.S.; SINEL'NIKOV, A.S.

Problem of efficient processing organization for shale retorts.
Trudy VNIIPS no.6:39-50 '58. (MIRA 11:8)
(Gas retorts) (Oil shales--Refining)

BARSHCHEVSKIY, M.M.; BEZMOZGIN, E.S.; ZAGLODIN, L.Z.; SINEL'NIKOV, A.S.;
SHUVALOV, V.I.

High production oil-shale retorts. Gaz. prom. no. 7:7-11 J1 '58.
(MIRA 11:7)

(Oil shales)
(Gas retorts)

BARS Hettensky, M.M.

PHASE I BOOK EXPLOITATION (CONT.)

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| | Mnichya i tekhnologiya topliv i produktov poerabotki, tpp. 9 (Chemistry and Technology of Fuel and Products of Refining), N.P. S. Lavrent'ev, Gontotekhnika Ltd., 1959, Ed. 2, (Series: 108: Leningrad, Gontotekhnika Ltd., 2,500 copies printed, frandy, Krafts'kaii Izdatel'stvo). | 9 |
| | Sponsoring Agency: R.S.F.S.R. Leninskej ekonomicheskoj Administrativnoj rayon. Sovet narodogo khozyaistva. | |
| | Ed.: V.M. Krush; Red.: A.A. Chishov; Tech. Ed.: A.B. Fashchentchikova; Editorial Board of Series: Z.G. Berestin, A.E. Drubin, D.K. Kollerov, S.S. Semenov, A.J. Snel'dinskij, and A.S. Poteyev. | |
| | PURPOSE: This collection of articles is intended for scientific, engineering and technical personnel in plants of the fuel and gas industry. | 632 |
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BARSHCHEVSKIY, M.M.; BEZMOZGIN, E.S.; SINEL'NIKOV, A.S.; SHUVALOV, V.I.

Shale-gas producers with a central feed for the heat-carrying
agent. Trudy VNIIPS no.7:120-146 '59. (MIRA 12:9)
(Gas retorts)

BEZMOZGIN, E.S.; BARSHCHEVSKIY, M.M.; VASIL'YEVA, M.M.

Prospects for the use of oxygen in the shale-gas industry. Trudy
VNIIT no.8:66-74 '59. (MIRA 13:4)
(Oil shales) (Oxygen)

BEZMOZGIN, E.S.; BARSHCHEVSKIY, M.M.; SINEL'NIKOV, A.S.; ZAGLODIN, L.S.
Increasing the capacity of pinch-type shale gas producers used at
the combine in Kohtla-Jarve. Trudy VNIIT no. 9:27-30 '60.
(MIRA 13:11)
(Kohtla-Jarve--Gas producers)

BARSHCHEVSKIY, M.M.; BEZMOZGIN, E.S.; VASIL'YEVA, M.M.; ZAGLODIN, L.S.;
SINEL'NIKOV, A.S.
Efficient system of processing Baltic oil shales. Trudy VNIIT no.9:
4-9 '60. (Oil shales) (MIRA 13:11)

BEZMOZGIN, E.S., BARSHCHEVSKIY, M.M., SINEL'NIKOV, A.S., ZAGLODIN, L.S.
Industrial experience in increasing the capacity of oil shale gas
producers. Gaz.prom. 5 no.2:17-19 F '60. (MIRA 13:6)
(Gas producers) (Oil shales)

BARSHCHEVSKIY, M.M.; BEZMOZGIN, E.S.; NEMCHENKO, A.G.; SINEL'NIKOV, A.S.

Experimental-plant testing of a newly designed reactor for the continuous
thermal-contact gasification of liquid fuel. Gaz.prom. 5 no.10:12-
16 O '60. (MIRA 13:10)
(Liquid fuels) (Gas manufacture and works)

BARSHEVSKIY, M.M.; BEZMOZGIN, E.S.; SHABIRO, P.N.

Investigating the segregation of fuel and the distribution of
gas flows on a cold model of a highly productive shale gas
generator. Trudy VNIIT no. 11:63-72 '62. (MIRA 17:5)

BABIN, I. N.; BAKSHCHEVSKIY, M. M.; BEZHOGIN, E. S.; PETROV, V. N.

Converting natural and mixed gas for special heating installations.
Trudy VNIIIT no. 11:245-253 '62. (MIRA 17:5)

BARSHCHEVSKIY, Mark Moiseyevich; BEZMOZGIN, Emmanuil Samuilovich;
SHAPIRO, Roal'd Natanovich; SINEL'NIKOV, A.S., nauchnyy
red.; SEGAL', Z.G., ved. red.; YASHCHURZHINSKAYA, A.B.,
tekhn. red.

[Handbook on refining oil shales] Spravochnik po pererabotke
goriuchikh slantsev. [By] M.M. Barshchevskii, E.S. Bezmoygin, R.N.
Shapiro, Leningrad, Gostoptekhizdat, 1963. 238 p.

(Oil shales)

(MIRA 16:3)

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U.S. GOVERNMENT, WASHINGTON, D.C.

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U.S. GOVERNMENT, WASHINGTON, D.C.
BY JAMES R. FISHER, DEPUTY CHIEF, TELETYPE SECTION.

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YUDKEVICH, Yu.D.; KUCHEROV, A.I.; and others, etc., etc.

Investigating the internal counterrevolutionary plot of the
Trudy VNIK no. 13: 100-17. - 64.

APPROVED FOR RELEASE: 06/06/2000

CIA-RDP86-00513R000203710015-2"

1-21117-65 SNT(-)/PPA(4)/T PR-4 WE

ACCESSION NR: AP5007147

S/0286/65/000/003/0012/0012

AUTHOR: Baranchevskiy, M. M.; Bazargin, E. S.; Kurdyukov, O. I.; Nemchenko, A. G.
Yudkevich, Yu. D.

TITLE: A method for thermal dissociation of fuel. Class 10, No. 167812

SOURCE: Byulleten' izobretaniy i tovarnykh znakov, no. 9, 1965, 12

TOPIC TAGS: fuel thermal decomposition, infrared radiation

ABSTRACT: This Author's Certificate introduces a method for thermal decomposition of fuel. In order to intensify the process, the dissociation is done under infrared radiation.

ASSOCIATION: none

SUBMITTED: 04Dec63

ENCL: 00

SUB CODE: PP

NO REF Sov: 000

OTHER: 000

Card 1/1

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BARSHCHEVSKIY, S.V., inzhener; KUZNETSOV, R.M., inzhener.

Mechanical fixing of cable shoes and connecting sleeves. Vest.elektroprom.
18 no.6:20-23 Je '47. (MLRA 6:12)

1. Leningradskiy elektrotekhnicheskiy institut im. V.I.Ul'yanova (Lenina).
(Electric cables)

BARSHCHEVSKIY, S.V.

KITAYENKO, G.I., laureat Stalinskoy premii; BARSHCHEVSKIY, S.V., retsenzent; POL'SKAYA, R.G., tekhnicheskij redaktor; MUGOKHANGATAF, E.A., tekhnicheskij redaktor.

[Electrician's handbook] Spravochnik elektromontazhnika. Leningrad,
Gos. soiuznos nauchno-tekhn. izd-vo sudostroit. promyshl. Vol.3.
1954. 639 p.
(Electric engineering)

(MIRA 8:5)

BARSICHEVSKIY, V.

Glider-gyroplane. Voen.znan. 25 no.9:24 S '49.
(MIRA 12:12)
(Helicopters)

BARSHCHEVSKIY, YU, A.

PARSHCHEVSKIY, YU, A. --"Construction Solutions of High Strength Based on
Ground Unslaked Lime." Min. Higher Education USSR, Kiev, 1955. (Dissertation
for the Degree of Candidate in Technical Sciences)

SO: Knizhnaya Letopis', No.35, 1955

PHASE I BOOK EXPLOITATION 904

Barshchevskiy, Yuriy Averkiyevich, Candidate of Technical Sciences

Bystrotverdeyushchiye rastvory; informatsionnoye soobshcheniye (Quick-Setting Solutions; Information Bulletin) Kiyev, 1957. 20 p. No. of copies printed not given.

Sponsoring Agencies: Ukrainian SSR. Ministerstvo stroitel'stva. Tekhnicheskoye upravleniye. Otdel tekhnicheskoy informatsii and Vsesoyuznyy nauchno-issledovatel'skiy institut organizatsii i mekhanizatsii promyshlenno-zhilishchnogo stroitel'stva.

Resp. Ed.: Orlovskaya, S.Ya.

PURPOSE: This information bulletin gives the results of work by the sponsoring agency in finding materials for quick-setting concrete solutions in an effort to decrease the working cycle and the area required for mass production of large cinder blocks.

COVERAGE: The bulletin covers the investigation of clinker (portland) and other types of cement. A time-material index and applicable formulae for the preparation of quick-setting solutions are given.

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Quick-Setting Solutions (Cont.) 904

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TM/sfm
1-12-59

BARSHCHEVSKIY, Yu., kand.tekhn. nauk (Kiyev).

Mortars for laying brick blocks. Stroi. mat. 4 no.8:35 Ag '58.
(MIRA 11:9)
(Mortar) (Building blocks)

RS. 11. 1.

"Influence of the Temperature on the Spectral Distribution of the Sensitivity
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